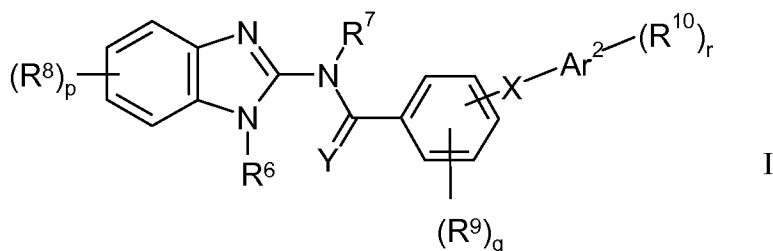


Listing of Claims:

1. (Currently amended) A compound or compounds of formula I



wherein

R^6, R^7 are independently selected from one another and are H, A or SO_2A , wherein, in the case of R^6 and R^7 , A is alkyl,

~~A is independently selected from the group consisting of alkyl, alkenyl, cycloalkyl, alkylencycloalkyl, alkoxy and alkoxyalkyl,~~

Ar^2 is phenyl, pyridinyl or pyrimidyl,~~selected independently from one another from aromatic hydrocarbons containing 6 to 14 carbon atoms and ethylenical unsaturated or aromatic heterocyclic residues containing 3 to 10 carbon atoms and one or two heteroatoms, independently selected from the group consisting of N, O and S,~~

R^{10} is selected from the group consisting of alkyl comprising 1 to 4 carbon atoms, alkoxy comprising 1 to 4 carbon atoms, Hal, CH_2Hal , $CH(Hal)_2$, perhaloalkyl comprising 1 to 4 carbon atoms, NO_2 , $(CH_2)_nCN$, $(CH_2)_nNR^{11}R^{12}$, $(CH_2)_nO(CH_2)_kNR^{11}R^{12}$, $(CH_2)_nCOR^{13}$, $(CH_2)_nCOOR^{13}$, $(CH_2)_nCONR^{11}R^{12}$,

$(\text{CH}_2)_n\text{SO}_2\text{NR}^{11}\text{R}^{12}$ and $(\text{CH}_2)_n\text{S}(\text{O})_u\text{R}^{13}$,

wherein

k is 0, 1 or 2,

r is 0, 1 or 2,

R^8 ; and R^9 ~~and~~ R^{10} are independently selected from the group consisting of

H, A, cycloalkyl comprising 3 to 7 carbon atoms, Hal, CH_2Hal , $\text{CH}(\text{Hal})_2$, $\text{C}(\text{Hal})_3$, NO_2 , and $(\text{CH}_2)_n\text{CN}$,

wherein, in the case of R^8 and R^{10} , A is independently selected

from the group consisting alkyl, alkenyl, cycloalkyl,

alkylenecycloalkyl, alkoxy and alkoxyalkyl,

$(\text{CH}_2)_n\text{NR}^{11}\text{R}^{12}$, $(\text{CH}_2)_n\text{OR}^{11}$, $(\text{CH}_2)_n\text{O}(\text{CH}_2)_k\text{NR}^{11}\text{R}^{12}$,

$(\text{CH}_2)_n\text{COOR}^{12}$, $(\text{CH}_2)_n\text{CONR}^{11}\text{R}^{12}$, $(\text{CH}_2)_n\text{NR}^{11}\text{COR}^{13}$,

$(\text{CH}_2)_n\text{NR}^{11}\text{CONR}^{11}\text{R}^{12}$, $(\text{CH}_2)_n\text{NR}^{11}\text{SO}_2\text{A}$, $(\text{CH}_2)_n\text{SO}_2\text{NR}^{11}\text{R}^{12}$,

$(\text{CH}_2)_n\text{S}(\text{O})_u\text{R}^{13}$, $(\text{CH}_2)_n\text{OC}(\text{O})\text{R}^{13}$, $(\text{CH}_2)_n\text{COR}^{13}$, $(\text{CH}_2)_n\text{SR}^{11}$,

$\text{CH}=\text{N}-\text{OA}$, $\text{CH}_2\text{CH}=\text{N}-\text{OA}$, $(\text{CH}_2)_n\text{NHOA}$, $(\text{CH}_2)_n\text{CH}=\text{N}-\text{R}^{11}$,

$(\text{CH}_2)_n\text{OC}(\text{O})\text{NR}^{11}\text{R}^{12}$, $(\text{CH}_2)_n\text{NR}^{11}\text{COOR}^{12}$,

$(\text{CH}_2)_n\text{N}(\text{R}^{11})\text{CH}_2\text{CH}_2\text{OR}^{13}$, $(\text{CH}_2)_n\text{N}(\text{R}^{11})\text{CH}_2\text{CH}_2\text{OCF}_3$,

$(\text{CH}_2)_n\text{N}(\text{R}^{11})\text{C}(\text{R}^{13})\text{HCOOR}^{12}$, $\text{C}(\text{R}^{13})\text{HCOOR}^{12}$,

$(\text{CH}_2)_n\text{N}(\text{R}^{11})\text{CH}_2\text{CH}_2\text{N}(\text{R}^{12})\text{CH}_2\text{COOR}^{12}$,

$(\text{CH}_2)_n\text{N}(\text{R}^{11})\text{CH}_2\text{CH}_2\text{NR}^{11}\text{R}^{12}$, $\text{CH}=\text{CHCOOR}^{11}$,

$\text{CH}=\text{CHCH}_2\text{NR}^{11}\text{R}^{12}$, $\text{CH}=\text{CHCH}_2\text{NR}^{11}\text{R}^{12}$, $\text{CH}=\text{CHCH}_2\text{OR}^{13}$,

$(\text{CH}_2)_n\text{N}(\text{COOR}^{11})\text{COOR}^{12}$, $(\text{CH}_2)_n\text{N}(\text{CONH}_2)\text{COOR}^{11}$,

$(\text{CH}_2)_n\text{N}(\text{CONH}_2)\text{CONH}_2$, $(\text{CH}_2)_n\text{N}(\text{CH}_2\text{COOR}^{11})\text{COOR}^{12}$,

$(\text{CH}_2)_n\text{N}(\text{CH}_2\text{CONH}_2)\text{COOR}^{11}$, $(\text{CH}_2)_n\text{N}(\text{CH}_2\text{CONH}_2)\text{CONH}_2$,

$(\text{CH}_2)_n\text{CHR}^{13}\text{COR}^{11}$, $(\text{CH}_2)_n\text{CHR}^{13}\text{COOR}^{11}$,

$(\text{CH}_2)_n\text{CHR}^{13}\text{CH}_2\text{OR}^{14}$, $(\text{CH}_2)_n\text{OCN}$ and $(\text{CH}_2)_n\text{NCO}$, wherein

A is independently selected from the group consisting of alkyl,

alkenyl, cycloalkyl, alkylenecycloalkyl, alkoxy and alkoxyalkyl,

- R^{11} , R^{12} are independently selected from the group consisting of H, A, $(CH_2)_mAr^3$ and $(CH_2)_mHet$, or in $NR^{11}R^{12}$,
- R^{11} and R^{12} form, together with the N-atom they are bound to, a 5-, 6- or 7-membered heterocycle which optionally contains 1 or 2 additional heteroatoms, selected from the group consisting of N, O and S,
- R^{13} , R^{14} ~~are~~ is independently selected from the group consisting of H, Hal, A, $(CH_2)_mAr^4$ and $(CH_2)_mHet$,
- Ar^3 , Ar^4 are independently selected from one another and are aromatic hydrocarbon residues comprising 5 to 12 carbon atoms which are optionally substituted by one or more substituents, selected from the group consisting of A, Hal, NO_2 , CN, OR^{15} , $NR^{15}R^{16}$, $COOR^{15}$, $CONR^{15}R^{16}$, $NR^{15}COR^{16}$, $NR^{15}CONR^{15}R^{16}$, $NR^{16}SO_2A$, COR^{15} , $SO_2R^{15}R^{16}$, $S(O)_uA$ and $OOCR^{15}$,
- Het is a saturated, unsaturated or aromatic heterocyclic residue which is optionally substituted by one or more substituents, selected from the group consisting of A, Hal, NO_2 , CN, OR^{15} , $NR^{15}R^{16}$, $COOR^{15}$, $CONR^{15}R^{16}$, $NR^{15}COR^{16}$, $NR^{15}CONR^{15}R^{16}$, $NR^{16}SO_2A$, COR^{15} , $SO_2R^{15}R^{16}$, $S(O)_uA$ and $OOCR^{15}$,
- R^{15} , R^{16} are independently selected from the group consisting of H, A, and $(CH_2)_mAr^6$, wherein
- Ar^6 is a 5- or 6-membered aromatic hydrocarbon which is optionally substituted by one or more substituents selected from the group

consisting of methyl, ethyl, propyl, 2-propyl, tert.-butyl, Hal, CN, OH, NH₂ and CF₃,

k, m and n are independently selected from ~~of~~ one another and are 0, 1, 2, 3, 4, or 5,

X is selected from the group consisting of O, S, CH₂, CH₂CH₂, OCH₂, CH₂O, C=O, C(=O)-NH and NH-C(=O) represents a bond or is (CR¹¹R¹²)_h, or (CHR¹¹)_h-Q-(CHR¹²)_i, wherein

~~Q ————— is selected from the group consisting of O, S, NR¹⁵, (CHal₂)_j, (O-CHR¹⁸)_j, (CHR¹⁸-O)_j, CR¹⁸=CR¹⁹, (O-CHR¹⁸CHR¹⁹)_j, (CHR¹⁸CHR¹⁹-O)_j, C=O, C=S, C=NR¹⁵, CH(OR¹⁵), C(OR¹⁵)(OR²⁰), C(=O)O, OC(=O), OC(=O)O, C(=O)N(R¹⁵), N(R¹⁵)C(=O), OC(=O)N(R¹⁵), N(R¹⁵)C(=O)O, CH=N-O, CH=N-NR¹⁵, S=O, SO₂, SO₂NR¹⁵ and NR¹⁵SO₂, wherein~~

~~R¹⁸, R¹⁹, R²⁰ ————— are independently selected from the meanings given for R⁸, R⁹ and R¹⁰;~~

~~h, i ————— are independently from each other 0, 1, 2, 3, 4, 5, or 6, and~~

~~j ————— is 1, 2, 3, 4, 5, or 6;~~

Y is selected from the group consisting of O, S, NR²¹, C(R²²)-NO₂, C(R²²)-CN and C(CN)₂, wherein

R²¹ ~~is independently selected from~~ has the meanings given for R¹³, R¹⁴ ~~and~~

R^{22} ~~is independently selected from~~ has the meanings given for R^{11} ,
or R^{12} ,

p, r ~~are~~ is independently from one another 0, 1, 2, 3, 4 or 5,

q is 0, 1, 2, 3 or 4,

u is 0, 1, 2 or 3,

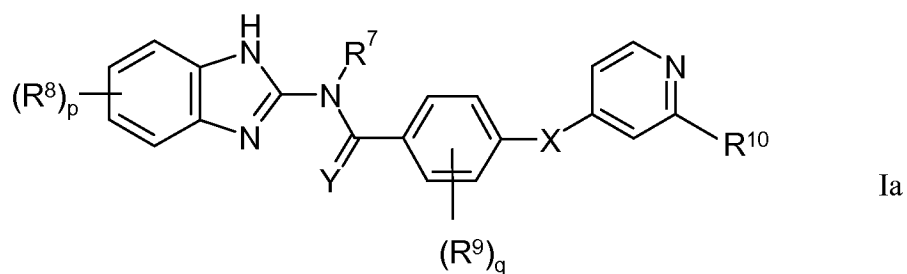
and

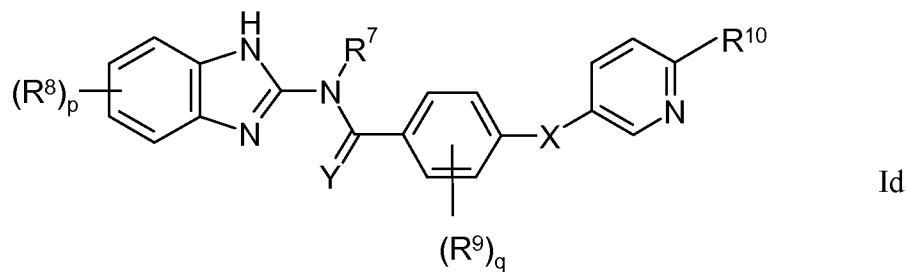
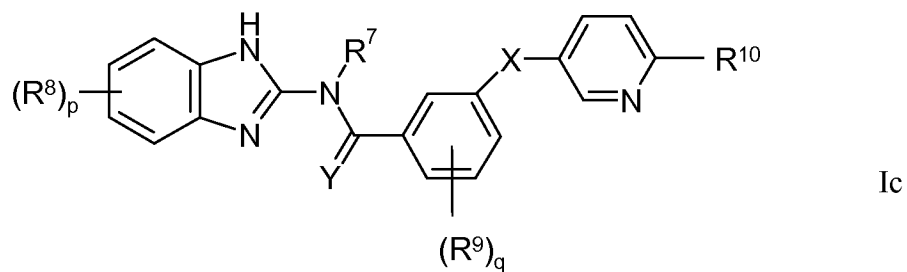
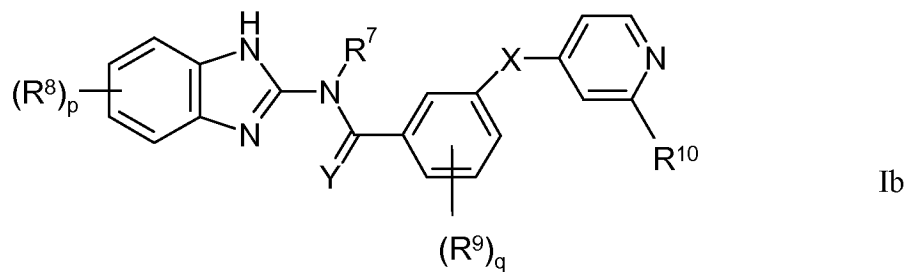
Hal is independently selected from the group consisting of F, Cl, Br
and I; or

physiologically acceptable derivatives, salts, solvates thereof or mixtures
thereof in all ratios.

2. (Canceled)

3. (Currently amended) The compound or compounds according to claim 1,
selected from the group consisting of the compounds of formulae Ia, Ib, Ic and
Id,





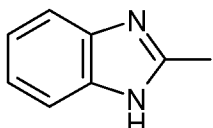
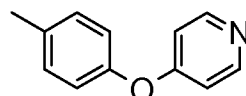
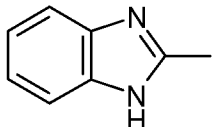
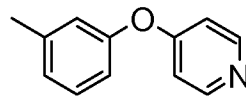
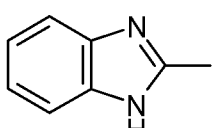
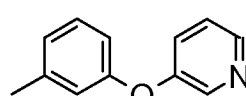
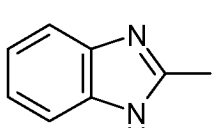
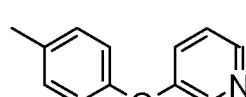
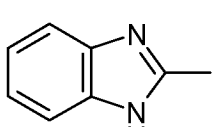
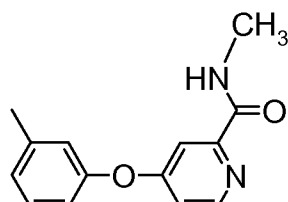
wherein

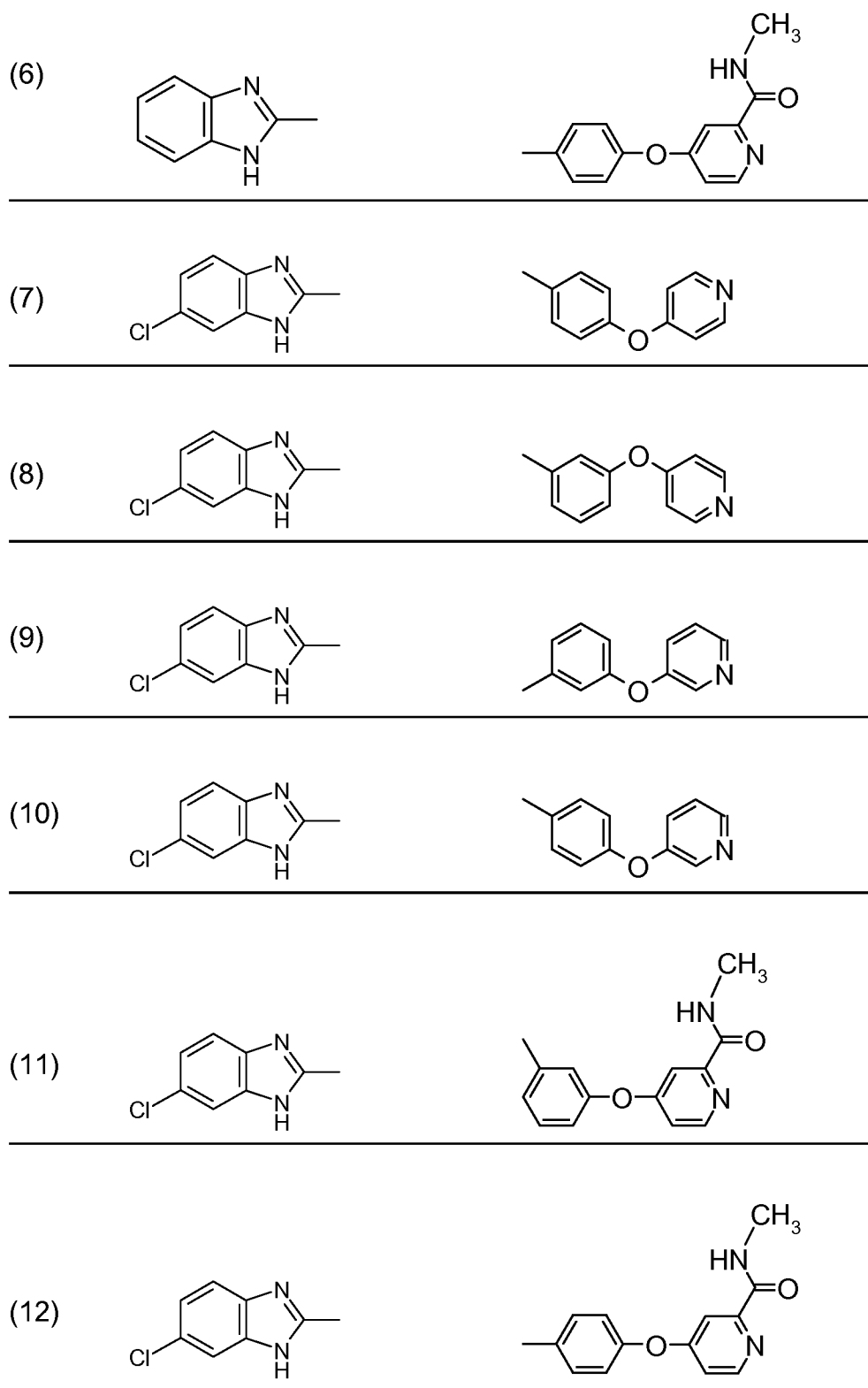
R^7 , R^8 , p , X , Y , R^9 and q are as defined in claim 1, and R^{10} is H or as defined in claim 1,

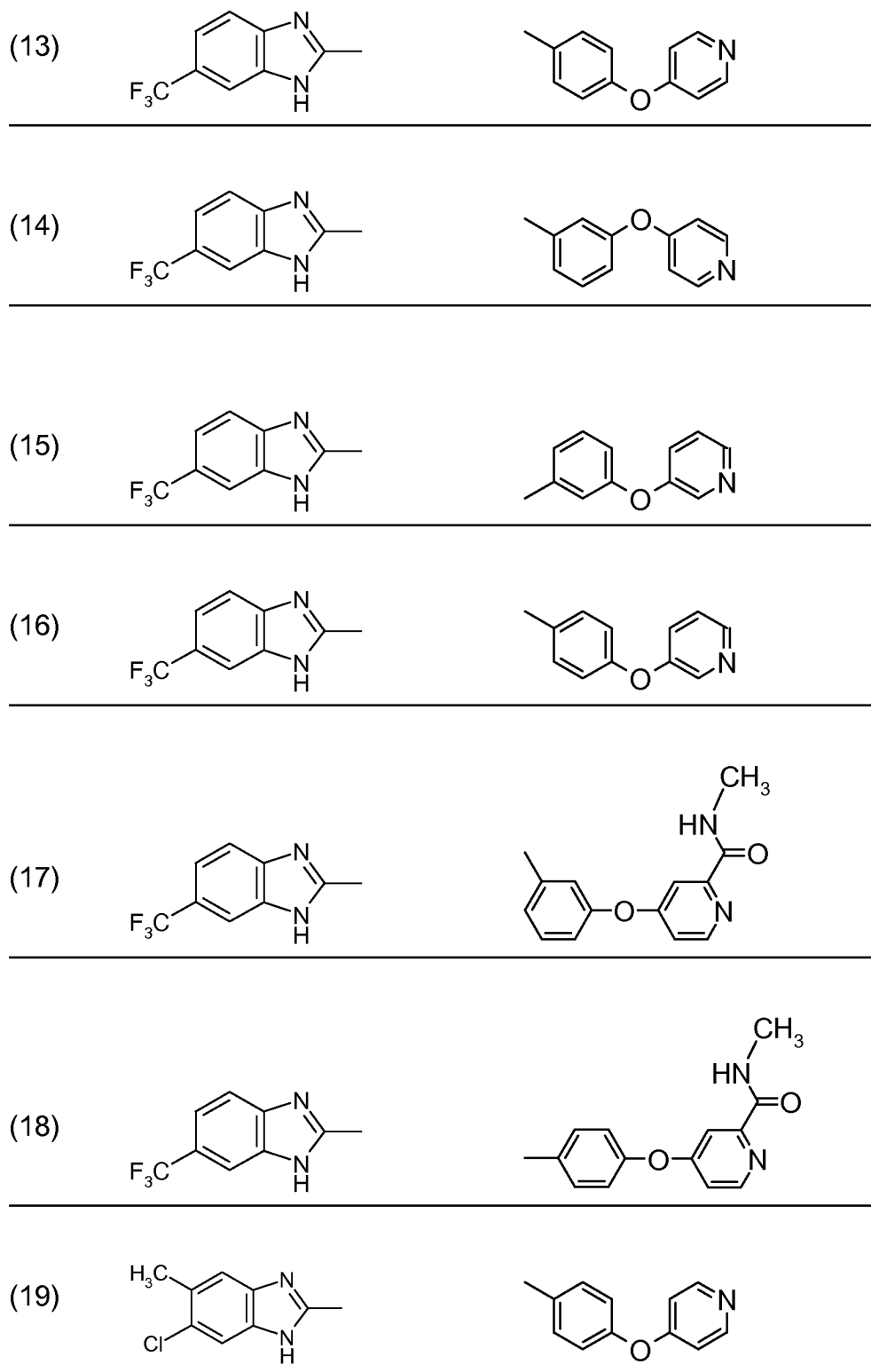
physiologically acceptable derivatives, salts, solvates thereof and mixtures thereof in all ratios.

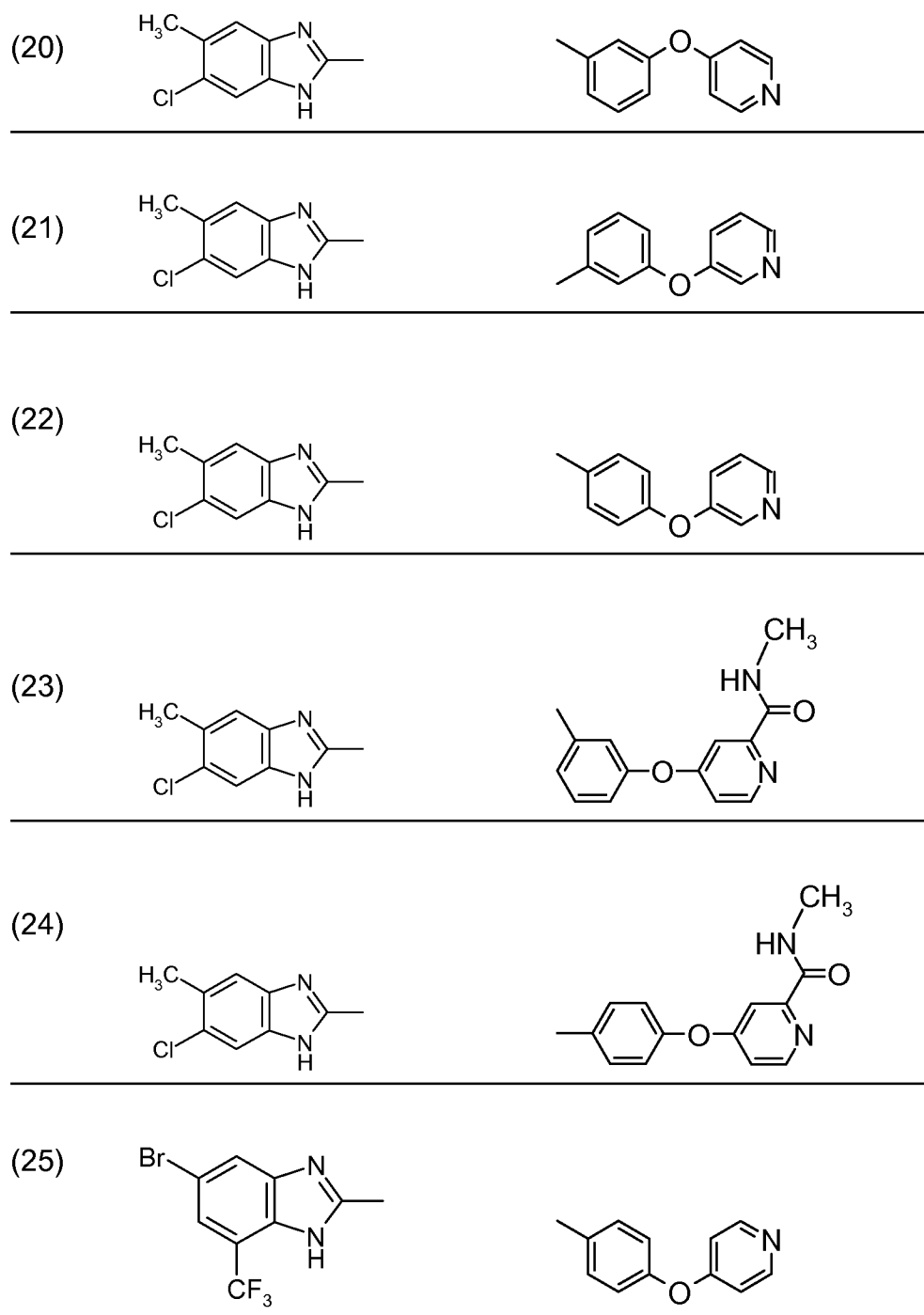
4. (Canceled)
5. (Currently amended) The compound or compounds according to claim 1, or tautomeric forms, pharmaceutically acceptable derivatives, solvates, salts,

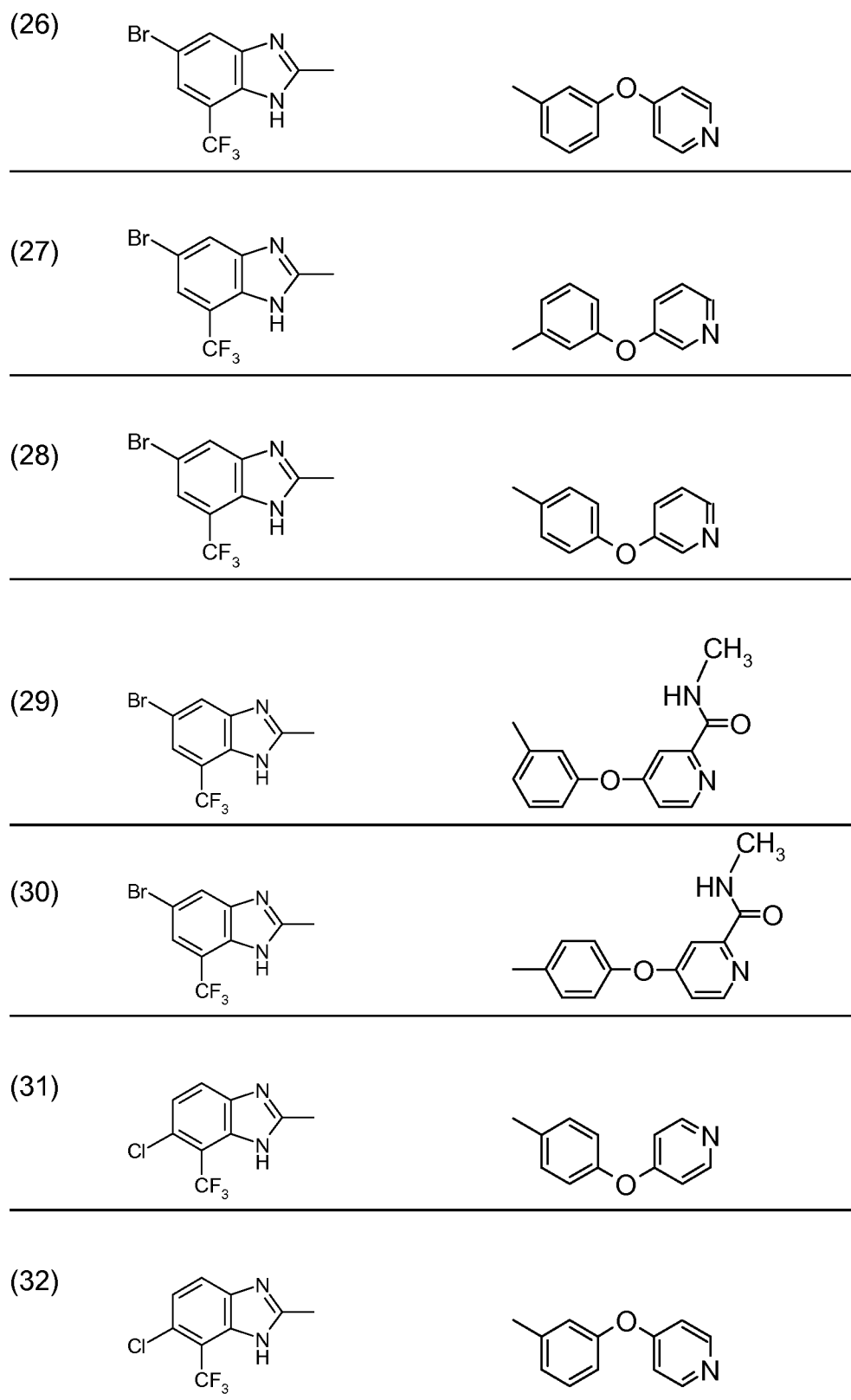
~~stereoisomers or mixtures thereof in all ratios, selected from the group consisting of compounds (1) to (78) of table 1; having formula A-CO-NH-B, wherein A- and -B are selected from the group consisting of, physiologically acceptable derivatives, salts, solvates, stereoisomers and mixtures thereof in all ratios~~

	<u>A-</u>	<u>-B</u>
(1)		
(2)		
(3)		
(4)		
(5)		

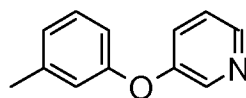
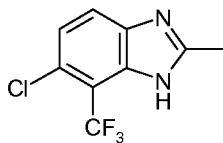




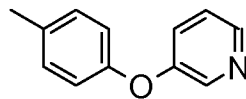
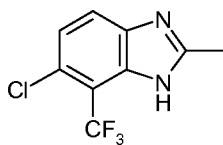




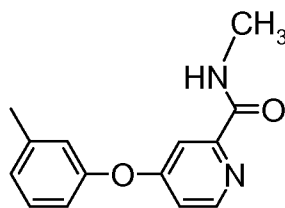
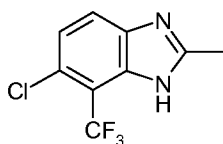
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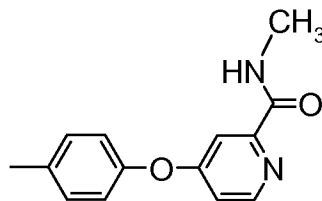
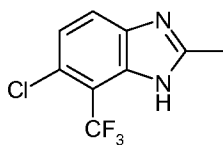
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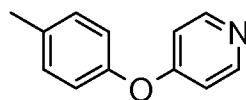
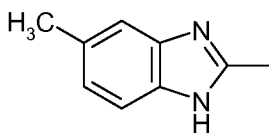
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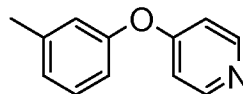
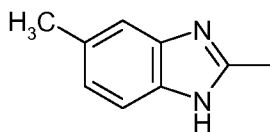
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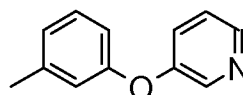
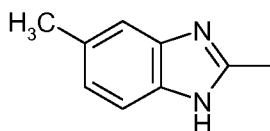
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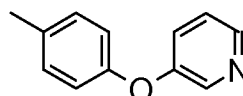
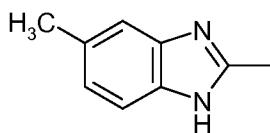
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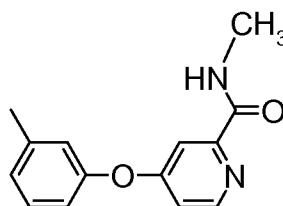
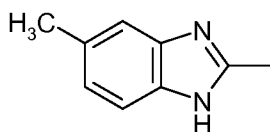
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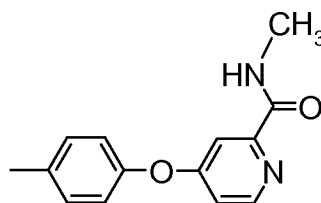
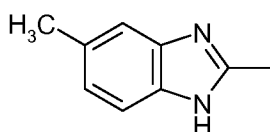
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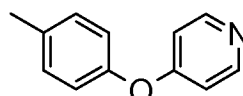
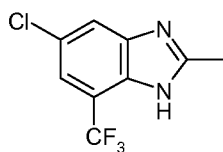
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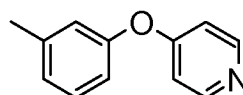
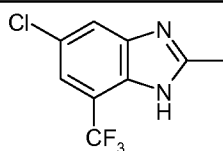
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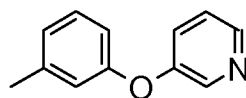
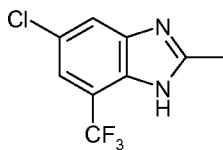
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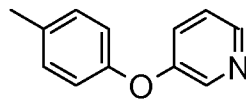
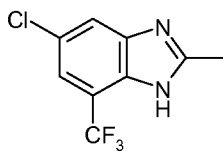
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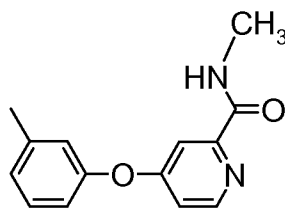
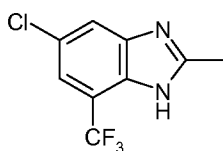
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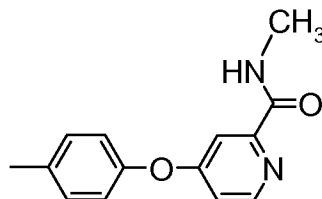
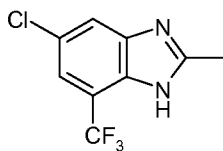
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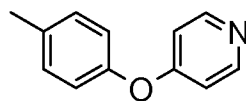
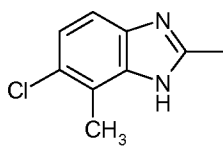
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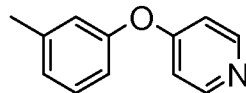
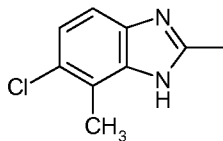
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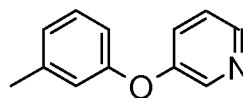
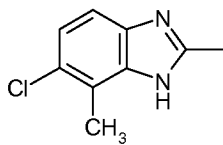
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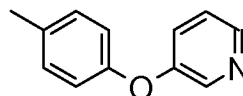
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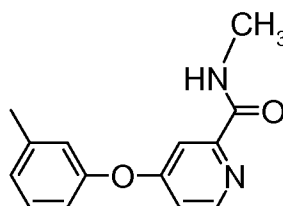
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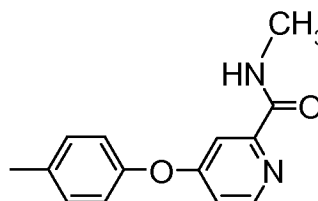
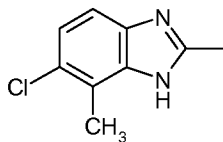
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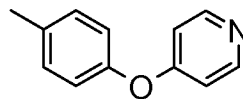
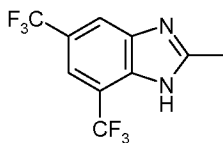
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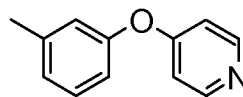
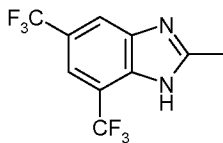
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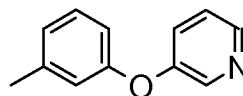
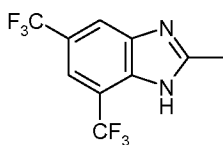
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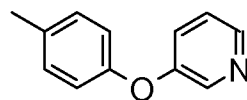
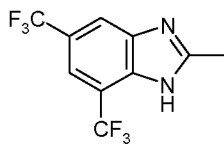
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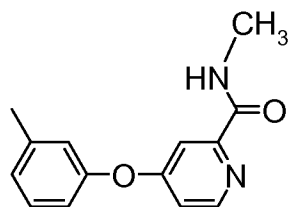
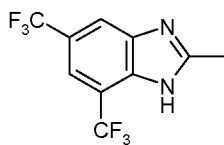
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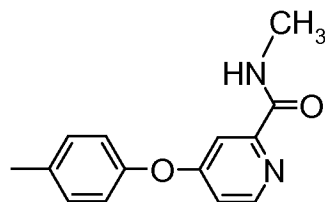
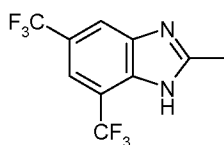
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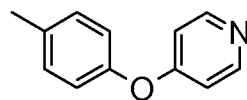
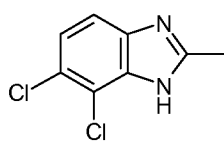
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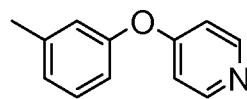
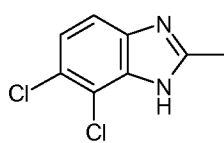
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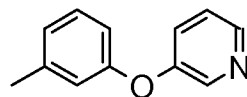
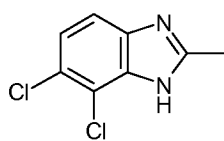
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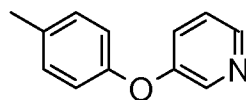
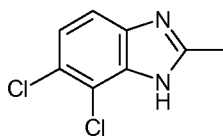
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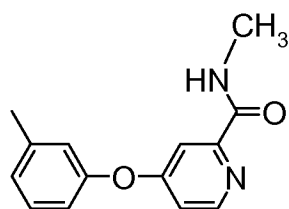
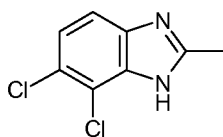
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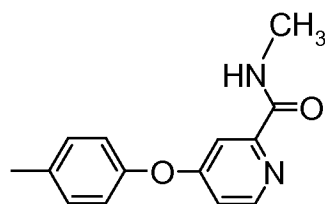
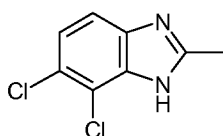
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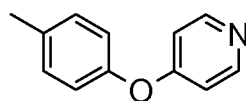
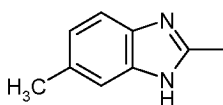
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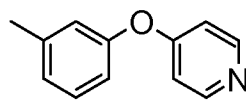
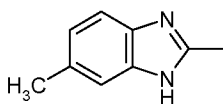
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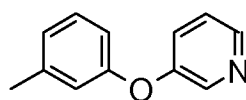
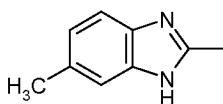
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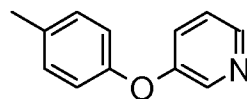
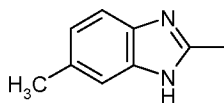
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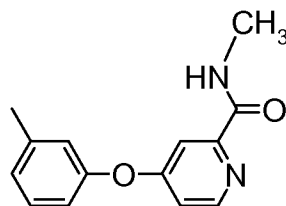
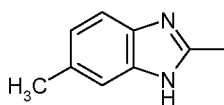
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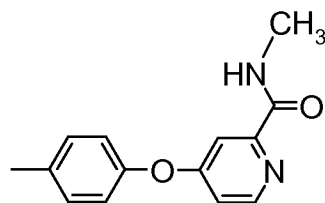
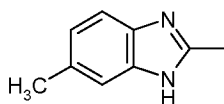
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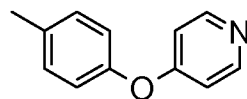
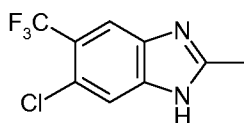
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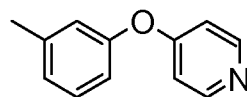
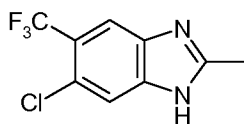
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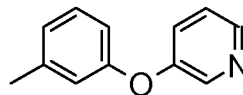
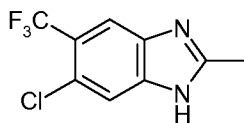
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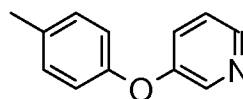
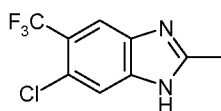
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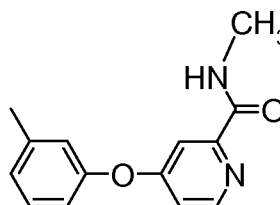
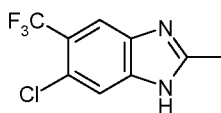
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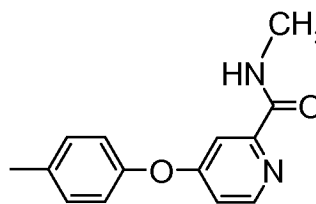
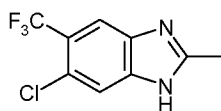
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(77)



(78)



6. (Currently amended) The compound or compounds according to claim 1, or tautomeric forms, pharmaceutically acceptable derivatives, solvates, salts, stereoisomers or mixtures thereof in all ratios, as a medicament.
7. (Currently amended) The compound or compounds according to claim 1, or tautomeric forms, pharmaceutically acceptable derivatives, solvates, salts, stereoisomers or mixtures thereof in all ratios, as a kinase inhibitor.

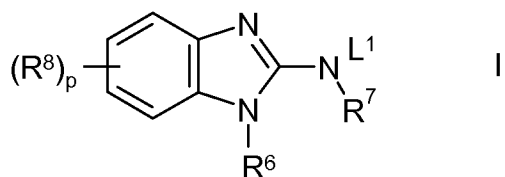
8. (Previous presented) The compound or compounds according to claim 7, characterized in that the kinases are selected from the group consisting of raf-kinases and VEGFR kinases.
9. (Currently amended) A pharmaceutical composition, comprising one or more of the compound or compounds according to claim 1, pharmaceutically acceptable derivatives, solvates, salts, stereoisomers or mixtures thereof in all ratios, as a pharmaceutical composition.
10. (Previously presented) The pharmaceutical composition according to claim 9, characterized in that it contains one or more additional compounds, selected from the group consisting of physiologically acceptable excipients, auxiliaries, adjuvants, carriers and pharmaceutical active ingredients.
11. (Currently amended) A process for the manufacture of a pharmaceutical composition, comprising one or more of the compound or compounds according to claim 1, or tautomeric forms, pharmaceutically acceptable derivatives, solvates, salts, stereoisomers or mixtures thereof in all ratios, and one or more compound or compounds, selected from the group consisting of carriers, excipients, auxiliaries and pharmaceutical active ingredients other than the compound or compounds according to claim 1, is processed by mechanical means into a pharmaceutical composition that is suitable as dosage form for application and/or administration to a patient.
12. (Withdrawn, currently amended) A method comprising administering to a patient the compound or compounds according to claim 1, or tautomeric forms, pharmaceutically acceptable derivatives, solvates, salts, stereoisomers or mixtures thereof in all ratios, as a pharmaceutical composition.

13. (Withdrawn, Currently amended) A method comprising administering to a patient the compound or compounds according to claim 1, or tautomeric forms, pharmaceutically acceptable derivatives, solvates, salts, stereoisomers or mixtures thereof in all ratios, in the treatment and/or prophylaxis of a disorder or disorders.
14. (Cancelled)
15. (Withdrawn) The method of claim 13, characterized in that the disorder or disorders are caused, mediated and/or propagated by kinases selected from the group consisting of raf-kinases and VEGFR kinases.
16. (Withdrawn) The method of claim 13, characterized in that the disorder or disorders are selected from the group consisting of hyperproliferative and nonhyperproliferative disorders.
17. (Withdrawn) The method of claim 13, characterized in that the disorder or disorders is cancer.
18. (Withdrawn) The method of claim 13, characterized in that the disorder or disorders is noncancerous.
19. (Withdrawn) The method of claim 18, characterized in that the noncancerous disorder or disorders are selected from the group consisting of infection, psoriasis, arthritis, inflammation, endometriosis, scarring, benign prostatic hyperplasia, immunological disease, autoimmune disease and immunodeficiency disease.
20. (Withdrawn, currently amended) The method of claim 17, characterized in that the cancer is ~~disorder or disorders are~~ selected from the group consisting of

brain cancer, lung cancer, squamous cell cancer, bladder cancer, gastric cancer, pancreatic cancer, hepatic cancer, renal cancer, colorectal cancer, breast cancer, head cancer, neck cancer, oesophageal cancer, gynaecological cancer, thyroid cancer, lymphoma, chronic leukaemia and acute leukaemia.

21. (Withdrawn) The method of claim 13 characterized in that the disorder or disorders are selected from the group consisting of arthritis, restenosis; fibrotic disorders; mesangial cell proliferative disorders, diabetic nephropathy, malignant nephrosclerosis, thrombotic microangiopathy syndromes, organ transplant rejection, glomerulopathies, metabolic disorders, inflammation and neurodegenerative disease.
22. (Withdrawn) The method of claim 13, characterized in that the disorder or disorders are selected from the group consisting of rheumatoid arthritis, inflammation, autoimmune disease, chronic obstructive pulmonary disease, asthma, inflammatory bowel disease, fibrosis, atherosclerosis, restenosis, vascular disease, cardiovascular disease, inflammation, renal disease and angiogenesis disorders.
23. (Withdrawn, currently amended) A method of treatment comprising administering to a patient the compound or compounds according to claim 1, or tautomeric forms, pharmaceutically acceptable derivatives, solvates, salts, stereoisomers or mixtures thereof in all ratios, as a kinase inhibitor.
24. (Withdrawn) The method of claim 23, characterized in that the kinase is one or more raf-kinases, selected from the group consisting of A-Raf, B-Raf and Raf-1.
25. (Canceled)

26. (Canceled)
27. (Canceled)
28. (Withdrawn) The method of claim 17, characterized in that the disorder or disorders is cancerous cell growth mediated by one or more kinases.
29. (Withdrawn, currently amended) A method for producing the compound or compounds of claim 1, or tautomeric forms, pharmaceutically acceptable derivatives, solvates, salts, stereoisomers or mixtures thereof in all ratios, comprising that
- a) a compound of formula II

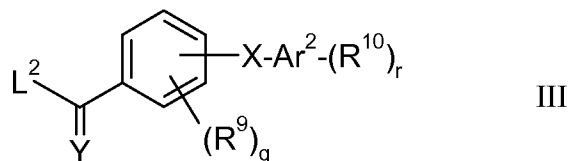


wherein

L^1 is H or a metal ion, and R^6 , R^7 , R^8 and p are as defined in claim 1,

is reacted

b) with a compound of formula III,



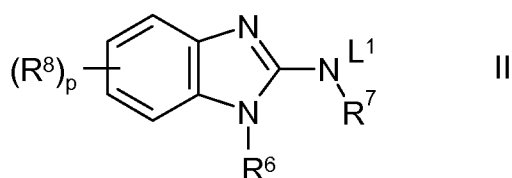
wherein

L^2 is Cl, Br, I, OH, an esterified OH-group or a diazonium moiety, and Y, R^9 , q , X, Ar^2 , R^{10} and r are as defined in claim 1,

and optionally

c) isolating and/or treating the compound or compounds of claim 1 obtained by said reaction with an acid, to obtain the salt thereof.

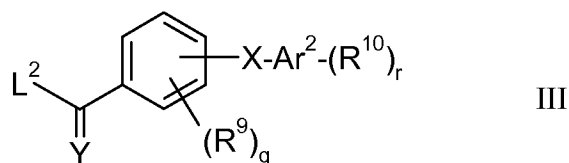
30. (Withdrawn) A compound or compounds of formula II,



wherein

L^1 is H or a metal ion, and R^6 , R^7 , R^8 and p are as defined in claim 1.

31. (Withdrawn) A compound or compounds of formula III,



wherein

L^2 is Cl, Br, I, OH, an esterified OH-group or a diazonium moiety, and Y, R^9 , q, X, Ar^2 , R^{10} and r are as defined in claim 1.

32. (New) The compound or compounds according to claim 1,

wherein

R^6 and R^7 are independently from one another H or alkyl, wherein alkyl is a unbranched or branched alkyl residue comprising 1 to 6 carbon atoms, optionally optionally substituted by one or more halogen atoms, by one or more hydroxy groups or by one or more amino groups which can optionally be substituted by alkyl comprising 1 to 6 carbon atoms,

Ar^2 is pyridinyl or pyrimidyl,

R^8 is independently selected from the group consisting of H, hal, unbranched or branched alkyl residues comprising 1 to 6 carbon atoms, optionally substituted by one or more halogen atoms, by one or more hydroxy groups or by one or more amino groups which can optionally be substituted by alkyl comprising 1 to 6 carbon atoms, and unbranched or branched alkoxy residues comprising 1 to 6 carbon atoms, optionally substituted by one or more halogen

atoms, by one or more hydroxy groups or by one or more amino groups which can optionally be substituted by alkyl comprising 1 to 6 carbon atoms,

R^9 is independently selected from the group consisting of H, hal, and unbranched or branched alkyl residues comprising 1 to 6 carbon atoms, optionally substituted by one or more halogen atoms, by one or more hydroxy groups or by one or more amino groups which can optionally be substituted by alkyl comprising 1 to 6 carbon atoms,

R^{10} is independently selected from the group consisting of H, alkyl comprising 1 to 4 carbon atoms, $(CH_2)_nNR^{11}R^{12}$, $(CH_2)_nO(CH_2)_kNR^{11}R^{12}$, $(CH_2)_nCOR^{13}$, $(CH_2)_nCOOR^{13}$, $(CH_2)_nCONR^{11}R^{12}$,

X is selected from the group consisting of O, S and CH_2 ,

Y is selected from the group consisting O, S and NR^{21} ,

or tautomeric forms, pharmaceutically acceptable derivatives, solvates, salts, stereoisomers or mixtures thereof in all ratios.

33. (New) The compound or compounds according to claim 1, wherein

R^6 and R^7 are independently from one another H or alkyl, wherein alkyl is selected from the group consisting of methyl, ethyl, trifluoro methyl, pentafluoro ethyl, isopropyl, tert.-butyl, 2-amino ethyl,

N-methyl-2-amino ethyl, N,N-dimethyl-2-amino ethyl,
N-ethyl-2-amino ethyl, N,N-diethyl-2-amino ethyl, 2-hydroxy
ethyl, 2-methoxy ethyl and 2-ethoxy ethyl,

Ar² is pyridinyl or pyrimidyl,

R⁸ is independently selected from the group consisting of H, hal,
alkyl residues selected from the group consisting of methyl,
ethyl, trifluoro methyl, pentafluoro ethyl, isopropyl, tert.-butyl,
2-amino ethyl, N-methyl-2-amino ethyl, N,N-dimethyl-2-amino
ethyl, N-ethyl-2-amino ethyl, N,N-diethyl-2-amino ethyl,
2-hydroxy ethyl, 2-methoxy ethyl and 2-ethoxy ethyl, and
alkoxy residues selected from the group consisting of methoxy,
ethoxy, n-propoxy, isopropoxy, 2-butoxy, tert.-butoxy and
perhalogenated derivatives thereof selected from the group
consisting of O-CCl₃, O-CF₃, O-C₂Cl₅, O-C₂F₅, O-C(CCl₃)₃ and
O-C(CF₃)₃,

R⁹ is independently selected from the group consisting of H, hal,
and alkyl, wherein alkyl is selected from the group consisting of
methyl, ethyl, trifluoro methyl, pentafluoro ethyl, isopropyl,
tert.-butyl, 2-amino ethyl, N-methyl-2-amino ethyl,
N,N-dimethyl-2-amino ethyl, N-ethyl-2-amino ethyl,
N,N-diethyl-2-amino ethyl, 2-hydroxy ethyl, 2-methoxy ethyl
and 2-ethoxy ethyl,

R¹⁰ is independently selected from the group consisting of H, alkyl
comprising 1 to 4 carbon atoms, (CH₂)_nNR¹¹R¹²,
(CH₂)_nO(CH₂)_kNR¹¹R¹², (CH₂)_nCOR¹³, (CH₂)_nCOOR¹³ and
(CH₂)_nCONR¹¹R¹², wherein

R^{11} , R^{12} and R^{13} are independently selected from a group consisting of H and alkyl, wherein alkyl is selected from the group consisting of methyl, ethyl, trifluoro methyl, pentafluoro ethyl, isopropyl, tert.-butyl, 2-amino ethyl, N-methyl-2-amino ethyl, N,N-dimethyl-2-amino ethyl, N-ethyl-2-amino ethyl, N,N-diethyl-2-amino ethyl, 2-hydroxy ethyl, 2-methoxy ethyl and 2-ethoxy ethyl,

hal is independantly selected from F, Cl and Br,

X is selected from the group consisting of O, S and CH_2 , and

Y is selected from the group consisting O and S,

or tautomeric forms, pharmaceutically acceptable derivatives, solvates, salts, stereoisomers or mixtures thereof in all ratios